

Class XII
Chemistry
Chapter 7:

The p-Block Elements

Top concepts:

1. **p-Block elements:** Elements belonging to groups 13 to 18 of the periodic table are called p-block elements.
2. **General electronic configuration of p-block elements:** The p-block elements are characterized by the ns^2np^{1-6} valence shell electronic configuration.
3. **Representative elements:** Elements belonging to the s and p-blocks in the periodic table are called the representative elements or main group elements.
4. **Inert pair effect:** The tendency of ns^2 electron pair to participate in bond formation decreases with the increase in atomic size. Within a group the higher oxidation state becomes less stable with respect to the lower oxidation state as the atomic number increases. This trend is called 'inert pair effect'. In other words, the energy required to unpair the electrons is more than energy released in the formation of two additional bonds.
5. **Nitrogen family:** The elements of group 15 – nitrogen (N), phosphorus (P), arsenic (As), antimony (Sb) and bismuth (Bi) belong to configuration is ns^2np^3 .
6. **Oxygen family:** Group 16 of periodic table consists of five elements – oxygen (O), sulphur (S), selenium (Se), tellurium (Te) and polonium (Po). Their general electronic configuration is ns^2np^4 .
7. **The halogen family:** Group 17 elements, fluorine (F), chlorine (Cl), bromine (Br), iodine (I) and astatine (At), belong to halogen family. Their general electronic configuration is ns^2np^5 .
8. **Group 18 elements:** Helium (He), neon (Ne), argon (Ar), krypton (Kr), xenon (Xe), and radon (Rn) are Group 18 elements. They are also called noble gases. Their general electronic configuration is ns^2np^6 except helium which has electronic configuration $1s^2$. They are called noble gases because they show very low chemical reactivity.

GROUP 15 ELEMENTS

9. **Atomic and ionic radii:** Covalent and ionic radii increase down the group. There is appreciable increase in covalent radii from N to P.

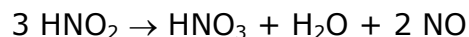
There is small increase from As to Bi due to presence of completely filled d or f orbitals in heavy elements.

10. **Ionisation energy:** It goes on decreasing down the group due to increase in atomic size. Group 15 elements have higher ionisation energy than group 14 elements due to smaller size of group 15 elements. Group 15 elements have higher ionization energy than group 16 elements because they have stable electronic configuration i.e., half filled p-orbitals.
11. **Allotropy:** All elements of Group 15 except nitrogen show allotropy.
12. **Catenation:** Nitrogen shows catenation to some extent due to triple bond but phosphorus shows catenation to maximum extent. The tendency to show catenation decreases down the group.
13. **Oxidation states:** The common oxidation states are +3, +5, -3. The tendency to show -3 oxidation state decreases down the group due to decrease in electronegativity which is due to increase in atomic size.

The stability of +5 oxidation state decreases whereas stability of +3 oxidation state increases due to inert pair effect.

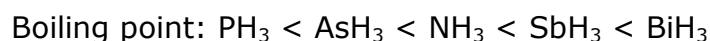
Nitrogen shows oxidation states from -3 to +5.

Nitrogen and phosphorus with oxidation states from +1 to +4 undergo oxidation as well as reduction in acidic medium. This process is called disproportionation.

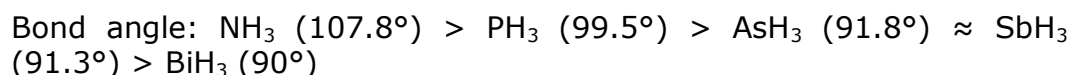


14. **Reactivity towards hydrogen:** All group 15 elements form trihydrides, MH_3 . Hybridisation - sp^3

The stability of hydrides decrease down the group due to decrease in bond dissociation energy down the group.



Boiling point increases with increase in size due to increase in van der Waals forces. Boiling point of NH_3 is more because of hydrogen bonding.



Electronegativity of N is highest. Therefore, the lone pairs will be towards nitrogen and hence more repulsion between bond pairs. Therefore bond angle is the highest. After nitrogen, the electronegativity decreases down the group.

Basicity decreases as $\text{NH}_3 > \text{PH}_3 > \text{AsH}_3 > \text{SbH}_3 < \text{BiH}_3$.

This is because the lone pair of electrons are concentrated more on nitrogen and hence the basicity will be maximum in the case of NH_3 . It will decrease down the group as the electronegativity decreases down the group.

The reducing power of hydrides increases down the group due to decrease in bond dissociation energy down the group.

15. **Reactivity towards oxygen:** All group 15 elements form trioxides (M_2O_3) and pentoxides (M_2O_5).

Acidic character of oxides decreases and basicity increases down the group. This is because the size of nitrogen is very small. It has a strong positive field in a very small area. Therefore, it attracts the electrons of water's O-H bond to itself and release H^+ ions easily. As we move down the group, the atomic size increases and hence the acidic character of oxides decreases and basicity increases down the group.

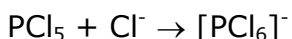
16. **Reactivity towards halogen:** Group 15 elements form trihalides and pentahalides.

Trihalides – covalent compounds and become ionic down the group.

sp^3 hybridisation, pyramidal shape

Pentahalides - sp^3d hybridisation, TBP shape

They are lewis acids because of the presence of vacant d – orbitals.



PCl_5 is ionic in solid state and exist as $[\text{PCl}_4]^+ [\text{PCl}_6]^-$

In PCl_5 , there are three equatorial bonds and two axial bonds. The axial bonds are longer than equatorial bonds because of greater repulsion from equatorial bonds.

Nitrogen does not form pentahalides due to absence of d- orbitals.

17. **Reactivity towards metals:** All elements react with metals to form binary compounds in -3 oxidation state.

18. **Anomalous behaviour of nitrogen:** The behaviour of nitrogen differs from rest of the elements.

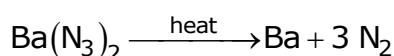
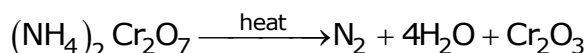
Reason:

- i. It has a small size.

- ii. It does not have d – orbitals
- iii. It has high electronegativity
- iv. It has high ionization enthalpy

19. **Dinitrogen:**

Preparation:



Properties:

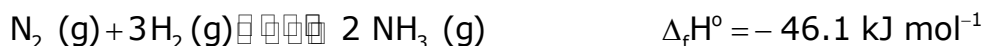
It is a colourless, odourless, tasteless and non – toxic gas. It is chemically un-reactive at ordinary temperature due to triple bond in

$\text{N} \equiv \text{N}$ which has high bond dissociation energy.

20. **Ammonia:** Ammonia molecule is trigonal pyramidal with nitrogen atom at the apex. It has 3 bond pairs and 1 lone pair. N is sp^3 hybridised.

Preparation:

Haber's process:



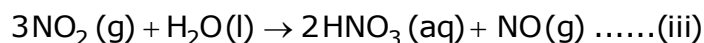
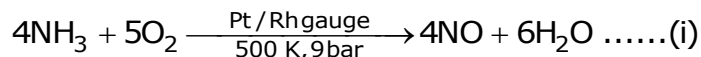
Pressure = $200 \times 10^5 \text{ Pa}$

Temperature = 773 K

Catalyst is FeO with small amounts of K_2O and Al_2O_3

21. **Nitric Acid:**

a. Ostwald Process:



NO thus formed is recycled and the aqueous HNO_3 can be concentrated by distillation upto $\sim 68\%$ by mass. Further concentration to 98% can be achieved by dehydration with concentrated H_2SO_4 .

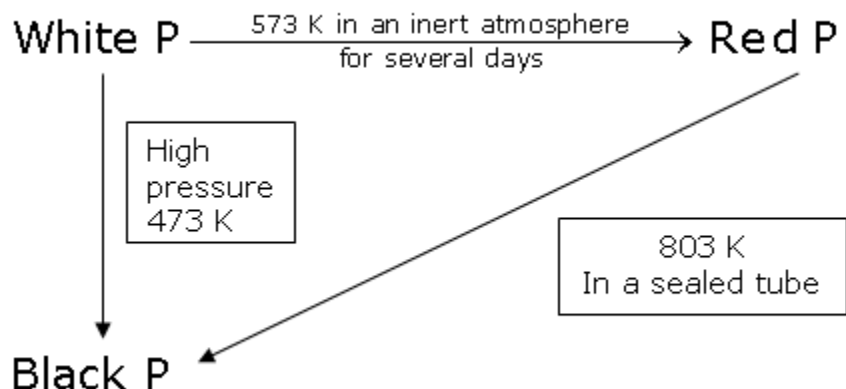
Nitric acid is strong oxidizing agent in the concentrated as well as in the dilute state.

22. Phosphorus:

- a. It shows the property of catenation to maximum extent due to most stable P – P bond.
- b. It has many allotropes, the important ones are :
 - i. White phosphorus
 - ii. Red phosphorus
 - iii. Black phosphorus

White phosphorus	Red phosphorus	Black phosphorus
Discrete tetrahedral P_4 molecules	Polymeric structure consisting of chains of P_4 units linked together	Exists in two forms - α black P and β black P
Very reactive	Less reactive than white P	Very less reactive
Glow in dark	Does not glow in dark	-
Translucent waxy solid	Has an iron grey luster	Has an opaque monoclinic or rhombohedral crystals
Soluble in CS_2 but insoluble in water	Insoluble in water as well as CS_2	-
It has low ignition temperature, therefore, kept under water	-	-

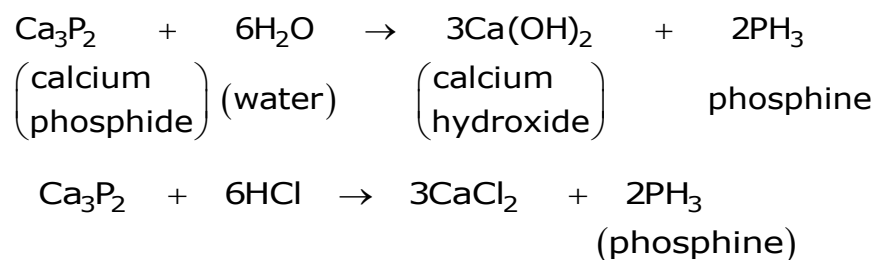
Preparation:



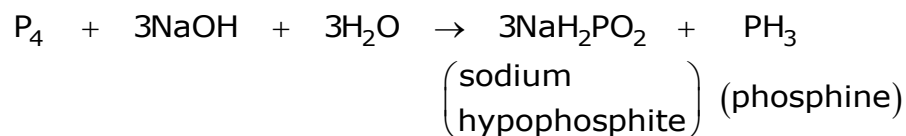
23. Phosphine:

Preparation:

i.



ii.



Phosphine is highly poisonous, colourless gas and has a smell of rotten fish.

24. Chlorides of Phosphorous:

PCl_3	PCl_5
Colourless oily liquid	Yellowish white powder

$P_4 + 6 Cl_2 \rightarrow 4 PCl_3$ $P_4 + 8 SOCl_2 \rightarrow 4 PCl_3 + 4 SO_2 + 2 S_2Cl_2$	$P_4 + 10 Cl_2 \rightarrow 4 PCl_5$ $P_4 + 10 SO_2Cl_2 \rightarrow 4 PCl_5 + 10 SO_2$
Is hydrolysed in the presence of moisture $PCl_3 + 3 H_2O \longrightarrow H_3PO_3 + 3 HCl$	$PCl_5 + H_2O \longrightarrow POCl_3 + 2 HCl$ $POCl_3 + 3 H_2O \longrightarrow H_3PO_4 + 3 HCl$
$3 CH_3COOH + PCl_3 \rightarrow 3 CH_3COCl + H_3PO_3$	$3 CH_3COOH + PCl_5 \rightarrow 3 CH_3COCl + POCl_3 + HCl$
$3 C_2H_5OH + PCl_3 \rightarrow 3 C_2H_5Cl + H_3PO_3$	$C_2H_5OH + PCl_5 \rightarrow C_2H_5Cl + POCl_3 + HCl$
Pyramidal shape, sp^3 hybridisation	TBP geometry, sp^3d hybridisation
-	$2 Ag + PCl_5 \rightarrow 2 AgCl + PCl_3$ $Sn + 2 PCl_5 \rightarrow SnCl_4 + 2 PCl_3$

GROUP 16 ELEMENTS

25. **Oxidation states:** They show -2, +2, +4, +6 oxidation states. Oxygen does not show +6 oxidation state due to absence of d - orbitals. Po does not show +6 oxidation state due to inert pair effect.

The stability of -2 oxidation state decreases down the group due to increase in atomic size and decrease in electronegativity.

Oxygen shows -2 oxidation state in general except in OF_2 and O_2F_2

The stability of +6 oxidation state decreases and +4 oxidation state increases due to inert pair effect.

26. **Ionisation enthalpy:** Ionisation enthalpy of elements of group 16 is lower than group 15 due to half filled p-orbitals in group 15 which are more stable. However, ionization enthalpy decreases down the group.

27. **Electron gain enthalpy:** Oxygen has less negative electron gain enthalpy than S because of small size of O.

From S to Po electron gain enthalpy becomes less negative to Po because of increase in atomic size.

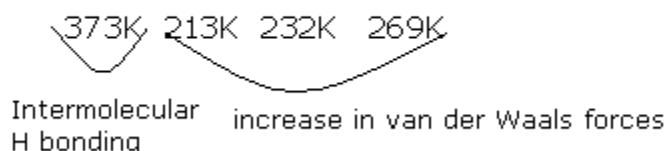
28. **Melting and boiling point:** It increases with increase in atomic number. Oxygen has much lower melting and boiling points than sulphur because oxygen is diatomic (O_2) and sulphur is octatomic (S_8).

29. **Reactivity with hydrogen:**

All group 16 elements form hydrides.

Bent shape

Bond angle: $H_2O > H_2S < H_2Se < H_2Te$



Acidic nature: $H_2O < H_2S < H_2Se < H_2Te$

This is because the H-E bond length increases down the group. Therefore, the bond dissociation enthalpy decreases down the group.

Thermal stability: $H_2O < H_2S < H_2Se < H_2Te < H_2Po$

This is because the H-E bond length increases down the group. Therefore, the bond dissociation enthalpy decreases down the group.

Reducing character: $H_2O < H_2S < H_2Se < H_2Te < H_2Po$

This is because the H-E bond length increases down the group. Therefore, the bond dissociation enthalpy decreases down the group.

30. **Reactivity with oxygen:** EO_2 and EO_3

Reducing character of dioxides decreases down the group because oxygen has a strong positive field which attracts the hydroxyl group and removal of H^+ becomes easy.

Acidity also decreases down the group.

SO_2 is a gas whereas SeO_2 is solid. This is because SeO_2 has a chain polymeric structure whereas SO_2 forms discrete units.

31. **Reactivity with halogens:** EX_2 EX_4 and EX_6

The stability of halides decreases in the order $F^- > Cl^- > Br^- > I^-$. This is because E-X bond length increases with increase in size.

Among hexa halides, fluorides are the most stable because of steric reasons.

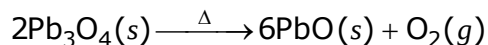
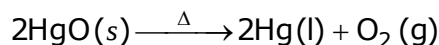
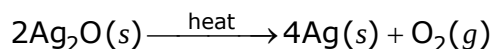
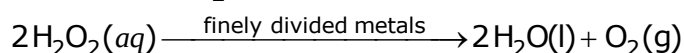
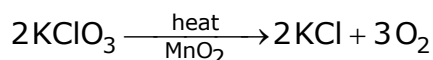
Dihalides are sp^3 hybridised, are tetrahedral in shape.

Hexafluorides are only stable halides which are gaseous and have sp^3d^2 hybridisation and octahedral structure.

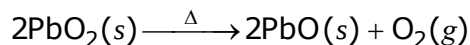
H_2O is a liquid while H_2S is a gas. This is because strong hydrogen bonding is present in water. This is due to small size and high electronegativity of O.

32. **Oxygen:**

Preparation:



(Red lead)



33. **Oxides:**

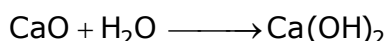
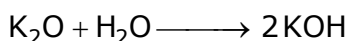
The compounds of oxygen and other elements are called oxides.

Types of oxides:

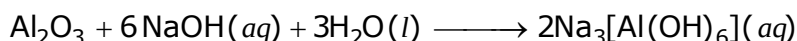
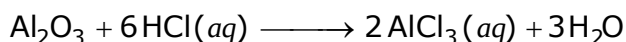
- a.** Acidic oxides: Non-metallic oxides are usually acidic in nature.



- b.** Basic oxides: Metallic oxides are mostly basic in nature. Basic oxides dissolve in water forming bases e.g.,



- c.** Amphoteric oxides: They show characteristics of both acidic as well as basic oxides.



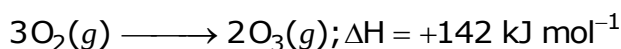
- d.** Neutral oxides: These oxides are neither acidic nor basic.

Example: Co, NO and N₂O

34. Ozone:

Preparation:

- i. It is prepared by passing silent electric discharge through pure and dry oxygen 10 – 15 % oxygen is converted to ozone.



Structure of Ozone: Ozone has angular structure. Both O = O bonds are of equal bond length due to resonance.

35. Sulphur:

Sulphur exhibits allotropy:

- Yellow Rhombic (α - sulphur):
- Monoclinic (β - sulphur):

α – Sulphur  ^{369 K}  β – Sulphur

At 369 K both forms are stable. It is called transition temperature.

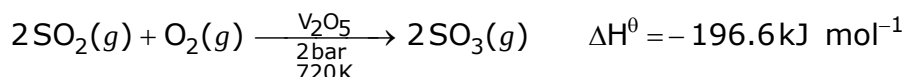
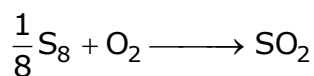
Both of them have S₈ molecules. The ring is puckered and has a crown shape.

Another allotrope of sulphur – cyclo S₆ ring adopts a chair form.

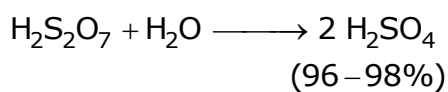
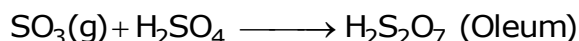
S₂ is formed at high temperature (~ 1000 K). It is paramagnetic because of 2 unpaired electrons present in anti bonding π^* orbitals like O₂

36. **Sulphuric acid:**

Preparation: By contact process



Exothermic reaction and therefore low temperature and high pressure are favourable



It is dibasic acid or diprotic acid.

It is a strong dehydrating agent.

It is a moderately strong oxidizing agent.

GROUP 17 ELEMENTS

37. **Atomic and ionic radii:** Halogens have the smallest atomic radii in their respective periods because of maximum effective nuclear charge.

38. **Ionisation enthalpy:** They have very high ionization enthalpy because of small size as compared to other groups.

39. **Electron gain enthalpy:** Halogens have maximum negative electron gain enthalpy because these elements have only one electron less than stable noble gas configuration.

Electron gain enthalpy becomes less negative down the group because atomic size increases down the group.

40. **Electronegativity:** These elements are highly electronegative and electronegativity decreases down the group. They have high effective nuclear charge.

41. **Bond dissociation enthalpy:**

Bond dissociation enthalpy follows the order $\text{Cl}_2 > \text{Br}_2 > \text{F}_2 > \text{I}_2$

This is because as the size increases bond length increases.

Bond dissociation enthalpy of Cl_2 is more than F_2 because there are large electronic repulsions of lone pairs present in F_2 .

42. **Colour:** All halogens are coloured because of absorption of radiations in visible region which results in the excitation of outer electrons to higher energy levels.

43. **Oxidising power:** All halogens are strong oxidising agents because they have a strong tendency to accept electrons.

Order of oxidizing power is $\text{F}_2 > \text{Cl}_2 > \text{Br}_2 > \text{I}_2$

44. **Reactivity with H_2 :**

Acidic strength: $\text{HF} < \text{HCl} < \text{HBr} < \text{HI}$

Stability: $\text{HF} > \text{HCl} > \text{HBr} > \text{HI}$

This is because of decrease in bond dissociation enthalpy.

Boiling point: $\text{HCl} < \text{HBr} < \text{HI} < \text{HF}$

HF has strong intermolecular H bonding

As the size increases van der Waals forces increases and hence boiling point increases.

% Ionic character: $\text{HF} > \text{HCl} > \text{HBr} > \text{HI}$

Dipole moment: $\text{HF} > \text{HCl} > \text{HBr} > \text{HI}$

Electronegativity decreases down the group.

Reducing power: $\text{HF} < \text{HCl} < \text{HBr} < \text{HI}$

45. **Reactivity with metals:** Halogens react with metals to form halides.

Ionic character: $\text{MF} > \text{MCl} > \text{MBr} > \text{MI}$

Halides in higher oxidation state will be more covalent than the one in the lower oxidation state.

46. **Interhalogen compounds: Reactivity of halogens towards other halogens:**

Binary compounds of two different halogen atoms of general formula

$\text{X X}'_n$ are called interhalogen compounds where $n = 1, 3, 5, \text{ or } 7$

These are covalent compounds.

All these are covalent compounds.

Interhalogen compounds are more reactive than halogens because X-X' is a more polar bond than X-X bond.

All are diamagnetic.

Their melting point is little higher than halogens.

XX' (ClF, BrF, BrCl, ICl, IBr, IF) (Linear shape)

XX'₃ (ClF₃, BrF₃, IF₃, ICl₃) (Bent T- shape)

XX'₅ – ClF₅, BrF₅, IF₅, (square pyramidal shape)

XX'₇ – IF₇ (Pentagonal bipyramidal shape)

47. **Oxoacids of halogens:**

Fluorine forms only one oxoacid HOF (Fluoric (I) acid or hypofluorous acid) due to high electronegativity.

Acid strength: HOCl < HClO₂ < HClO₃ < HClO₄

Reason: HClO₄ → H⁺ + ClO₄⁻

most stable

Acid strength: HOF > HOCl > HOBr > HOI

This is because Fluorine is most electronegative.

GROUP 18 ELEMENTS:

48. **Ionisation enthalpy:** They have very high ionization enthalpy because of completely filled orbitals.

Ionisation enthalpy decreases down the group because of increase in size.

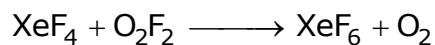
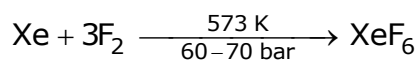
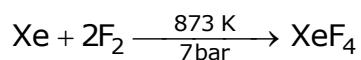
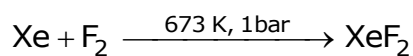
49. **Atomic radii:** Increases down the group because number of shells increases down the group.

50. **Electron gain enthalpy:** They have large electron gain enthalpy because of stable electronic configuration.

51. **Melting and boiling point:** Low melting and boiling point because only weak dispersion forces are present.

52. XeF₂ is linear, XeF₄ is square planar and XeF₆ is distorted octahedral. KrF₂ is known but no true compound of He Ne and Ar are known.

53. **Compounds of Xe and F:**



XeF_2 , XeF_4 and XeF_6 are powerful fluorinating agents.

54. **Compounds of Xe and O:**

